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Appropriation of GPIIb/IIIa from platelet-derived extracellular vesicles supports monocyte recruitment in systemic inflammation

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ABSTRACT

Interactions between platelets, leukocytes and the vessel wall provide alternative pathological routes of thrombo-inflammatory leukocyte recruitment. We found that when platelets were activated by a range of agonists in whole blood, they shed platelet-derived extracellular vesicles which rapidly and preferentially bound to blood monocytes compared to other leukocytes. Platelet-derived extracellular vesicle binding to monocytes was initiated by P-selectin-dependent adhesion and was stabilised by binding of phosphatidylserine. These interactions resulted in the progressive transfer of the platelet adhesion receptor GPIIb/IIIa to monocytes. GPIIb/IIIa⁺ monocytes tethered and rolled on immobilised von Willebrand Factor or were recruited and activated on endothelial cells treated with TGF-β1 to induce the expression of von Willebrand Factor. In both models monocyte adhesion was ablated by a function-blocking antibody against GPIIb/IIIa. Monocytes could also bind platelet-derived extracellular vesicle in mouse blood *in vitro* and *in vivo*. Intratracheal instillations of diesel nanoparticles, to model chronic pulmonary inflammation, induced accumulation of GPIIb/IIIa on circulating monocytes. In intravital experiments, GPIIb/IIIa⁺ monocytes adhered to the microcirculation of the TGF-β1-stimulated cremaster muscle, while in the *ApoE*^{-/-} model of atherosclerosis, GPIIb/IIIa⁺ monocytes adhered to the carotid arteries. In trauma patients, monocytes bore platelet markers within 1 hour of injury, the levels of which correlated with severity of trauma and resulted in monocyte clearance from the circulation. Thus, we have defined a novel thrombo-inflammatory pathway in which platelet-derived extracellular vesicles transfer a platelet adhesion receptor to monocytes, allowing their recruitment in large and small blood vessels, and which is likely to be pathogenic.

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Introduction

The recruitment of leukocytes during inflammation occurs in the haemodynamically permissive environment of the post capillary venules. In this environment, vascular endothelial cells responding to pro-inflammatory mediators such as cytokines express adhesion receptors and activating stimuli such as chemokines, which ensure efficient and localised trafficking of leukocytes into the affected tissues.^{1,4} It has become clear more recently that in pathological situations, platelets can also play a

role in leukocyte recruitment in other vascular beds.⁵ Thus, the integrated function of the thrombotic and inflammatory systems results in recruitment of leukocytes to arterioles in models of ischaemic injury of the liver and other tissues.⁶⁻¹⁰ Moreover, there is substantial evidence supporting a role for platelets in the preferential recruitment of monocytes to the artery wall during atherogenesis. For example, inhibition of platelet adhesion to the artery wall, or induction of thrombocytopenia, significantly reduces monocyte trafficking and the burden of atherosclerotic disease in genetically susceptible strains of mice.¹¹⁻¹⁴ In addition, instillation of activated platelets exacerbates the formation of atherosclerotic plaques in such models.¹¹⁻¹⁴ There is also direct evidence that platelet P-selectin plays a role in plaque formation in the *ApoE*^{-/-} mouse.¹¹⁻¹⁴ Other studies demonstrate that platelet derived chemokines such as CCL5 (RANTES) and CX3CL1 (fractalkine), once deposited on vascular endothelial cells, can selectively recruit monocytes in these models.¹¹⁻¹⁵

The examples described above require platelet activation at the vessel wall to facilitate leukocyte recruitment and trafficking. However, interactions between platelets and leukocytes also occur in circulating blood under pathological conditions. Indeed, formation of platelet-leukocyte aggregates has been described in diseases as diverse as bacterial infection, rheumatoid arthritis, diabetes and inflammatory bowel disease.¹⁶⁻²² In cardiovascular disease (CVD) the number of platelet-leukocyte aggregates increases significantly, and one can measure an increased incidence of such heterotypic aggregates in individuals with independent risk factors for CVD, such as hypertension.²³⁻²⁵ Indeed, it has been proposed that an increase in the incidence of platelet-leukocyte aggregates may in itself, be an independent risk factor for CVD.²⁶ The formation of platelet leukocyte aggregates may also play an important role in acute and severe inflammatory responses. Thus, in patients with acute trauma or trauma associated sepsis, an enhanced capacity for platelet activation and platelet interaction with monocytes and neutrophils has been reported in response to exogenous activation of their blood with the ionophore, ionomycin.^{27,28}

Extracellular vesicles which can be detected in the blood, urine and other bodily fluids are heterogeneous particles 40-1,500 nm in diameter that are derived from the plasma membrane (microvesicles) or by exocytosis of multi-vesicular bodies (exosomes).²⁹ They are released from cells of the vasculature, including platelets, endothelial cells (EC) and leukocytes, and specific populations can be identified using appropriate methodology (e.g. flow cytometry), as they express surface markers derived from their cell of origin. There is now mounting evidence that platelet-derived extracellular vesicles (PEV) (otherwise and often referred to as microparticles or microvesicles) are heterogeneous in nature. For example, *in vitro*, PEV have been generated in response to shear stress, thrombin, calcium ionophore, adenosine diphosphate (ADP), collagen and collagen related peptide.³⁰⁻³³ Interestingly, these studies show that PEV derive by using different platelet agonists and differ in abundance, as well as the cargo that they convey. Indeed, there is now good evidence that platelets can shed large vesicles which contain organelles such as mitochondria.³⁴ Until recent technological advancements it had been impossible to analyse the concentration and composition of vesicles using a single platform. Flow cytometry does not detect vesicles <200-

300 nm and does not accurately measure larger vesicles due to the disparity in the refractive index of biological vesicles and the latex beads used as size standards on this platform.³⁵ However, electron microscopy studies show that the majority of PEV are small. Thus, although Ponomereva *et al.* described calcium ionophore derived PEV as large as 1,500 nm, particles were predominantly in the range of 50-130 nm.³⁶ Similarly, Aatonen *et al.* described the main population of PEV as being 100-250 nm, with in excess of 90% of all vesicles being smaller than 500 nm irrespective of the platelet agonist used for PEV biogenesis.³⁵ Mitochondria containing vesicles, referred to above, were in the range of 500-1,500 nm. Importantly, the study of the functions of distinct subsets of PEV is not a well-developed field, however, bearing in mind the diversity of the PEV generated upon platelet activation, vesicles with discrete functional roles cannot be ruled out. The diversity of platelet microparticles has recently been reviewed.³⁷

There is mounting evidence that PEV play a pathophysiological role in inflammation.³⁸ An increased concentration of circulating PEV is associated with a number of diseases. In diabetic retinopathy, the number of PEV was associated with the severity of disease,³⁹ while the levels of PEV circulating in patients with type-1 diabetes correlated with the degree of pro-atherogenic dyslipidaemia.⁴⁰ There was a correlation with vascular dysfunction (assessed by measuring arterial elasticity and flow-dependent vasodilatation of the brachial artery) in patients with type-2 diabetes.⁴¹ Interestingly, the number of PEV was higher in patients with acute coronary syndromes than those with stable angina,⁴² implying an association with the onset of athero-thrombotic disease. The roles of PEV in inflammation and pathogenesis of inflammatory disease are not well understood. However, they possess adhesion receptors such as glycoprotein (GP)Ib α , α IIb β 3-integrin and P-selectin, meaning that they could interact with the vessel wall and circulating leukocytes to promote recruitment of the later. Importantly, as these receptors ordinarily regulate the process of haemostasis, PEV might provide an avenue of leukocyte recruitment to the disease environment which falls outside of regulatory pathways which ordinarily limit the duration and magnitude of the inflammatory response.

Here, using assays of heterotypic aggregate formation we have characterised the adhesive interactions between leukocytes and PEV in whole blood and identified a novel route by which the platelet adhesion receptor, GPIb α , promotes monocyte recruitment in both *in vitro* and *in vivo* models of vascular inflammation.

Methods

Full Methods can be found in the *Online Supplementary Materials and Methods*.

Blood donors

Blood was obtained from healthy donors with informed consent and with local ethical approval (ERN_07-058). Blood from the Golden Hour cohort (drawn within 1 hour of suffering traumatic injury) was obtained under the National Research Ethics Committee (reference 13/WA/0399). Specimen collection and informed consent procedures were approved and permission granted by the Biomedical Science Ethic Committee.

Animal experiments

All experiments were performed in accordance with the Home Office Guidelines. In each experiment C57BL/6 *IL4R/GPIb α -Tg* or *ApoE^{-/-}* or wild-type (WT) animals with the same background were allocated at random to experimental groups. Mice from the same litter were randomly distributed amongst experimental groups.

Results

Platelet activation in whole blood leads to formation of PEV and their adhesion to monocytes

We investigated the effect of platelet activation on platelet-leukocyte interactions in whole blood. When thrombin receptor activating peptide (TRAP), an agonist of the platelet protease activated receptor-1 (PAR-1), was added to sheared whole blood, a time dependent increase in the percentage of monocytes bearing the platelet receptor GPIb α (CD42b) as well as CD41 (GPIIb) and in the intensity of GPIb α and CD41 staining, was observed (Figure 1A-C; *Online Supplementary Figure S1A-C* and *S2*). In unstimulated blood, few monocytes (~5%) possessed measurable levels of GPIb α , showing that shear did not activate platelets. During analysis monocytes were subdivided into two subsets using standard markers as previously described.⁴³ Classical monocytes (CD14⁺CD16⁻) represent 90% of cells in the circulation and non-classical/intermediate monocytes (CD14⁺^{dim}CD16⁺) 10% (*Online Supplementary Figure S1A*). In our studies, we have compared classical to non-classical/intermediate monocytes grouped together. This is because the low numbers of isolated intermediate and non-classical monocytes do not allow appropriate functional testing of these subsets individually in our assays. The interaction between platelets and monocytes was similar when classical and non-classical/intermediate monocytes were assayed, showing similar patterns of GPIb α and CD41 accumulation over time (Figure 1B-C; *Online Supplementary Figure S1A-C* and *S2*). Interestingly, only a modest accumulation of GPIb α was evident on neutrophils stimulated with TRAP and even less when whole blood was stimulated with CRP-XL (Figure 1D and *Online Supplementary Figure S1D-E*). We observed no accumulation of GPIb α on lymphocytes (Figure 1D and *Online Supplementary Figure S1D*).

The median fluorescent intensity (MFI) of GPIb α on monocytes after 30 minutes (min) of TRAP stimulation was well below the intensity on individual platelets (Figure 2A). Moreover, the time course of the acquisition of GPIb α by monocytes demonstrated a progressive accumulation that ruled out the binding of whole platelets (Figure 2B). This pattern of accumulation is consistent with the adhesion of PEV, which was confirmed using confocal microscopy (Figure 2C and *Online Supplementary Figure S3*). For comparison, we show a monocyte bearing whole platelets generated under different experimental conditions *i.e.* in the absence of shear (Figure 2D).

Here, we have reported the formation of PEV in response to thromboxane A₂, ADP and cross linked collagen related peptide (CRP-XL). Activation of platelets in whole blood using CRP-XL, ADP, the thromboxane mimetic U46619, or the C-type lectin-like receptor (CLEC-2) agonist, rhodocytin, resulted in the same pattern of accumulation of GPIb α on monocytes, showing that different routes of platelet activation resulted in PEV pro-

duction and adhesion to monocytes (*Online Supplementary Figure S4A-D*).

PEV binding to monocytes is rapid

Accumulation of PEV on monocytes after stimulation of whole blood was progressive over a prolonged period of time (*i.e.* 30 min) (Figure 1B-C and *Online Supplementary Figure S4*). An important question is whether this pattern of accumulation is dependent upon the dynamics of PEV-monocyte interaction and adhesion, or whether the genesis of PEV from activated platelets is the rate-limiting step. Here we used the addition of isolated and pre-labelled PEV (1×10^9 /mL) generated by stimulating platelets (3×10^8) with CRP-XL ($1 \mu\text{g/mL}$), to unstimulated whole blood to investigate this. After CRP-XL activation of isolated platelets, GPIb α stained PEV were readily discernible by flow cytometry in platelet supernatants (Figure 3A and *Online Supplementary Figure S5A-E*). A similar pattern was observed for CD41 on PEV (*Online Supplementary Figure S5C* and *S6*). Interestingly, ~25% of the large vesicles detected by flow cytometry contained mitochondrial fragments, as previously described³⁴ (*Online Supplementary Figure S7A*). Analysis using nanoparticle tracking showed that 3×10^8 platelets could yield $1.2 \pm 0.3 \times 10^9$ PEV compared to an average $1.3 \times 10^8 \pm 2.8 \times 10^7$ vesicles in untreated conditions (Figure 3B and *Online Supplementary Figure S5D-E*) with a mean diameter of 274 ± 188 nm. To date it has not been possible to simultaneously count vesicles, size them and analyse protein cargo using a single platform. The Exo View-R100 is a new platform which allows such analysis providing previously unattainable information in a single protocol.⁴⁴ Using this assay we observed that PEV from CRP-XL stimulated platelets captured by a CD9 antibody had a mean size of 54 nm while those captured by a CD41a antibody had a mean size of 82.3 nm (*Online Supplementary Figure S7B*). Upon analysis using the Exo View system, we observed the majority of PEV were captured by CD41a and CD9 (which are abundant on platelets), but not CD63 or CD81 (which are expressed on exosomes) (*Online Supplementary Figure S7C*). This was also confirmed by secondary labelling of captured PEV using fluorescent antibodies against CD9, CD63 and CD81 (*Online Supplementary Figure S7D*). Labelled PEV were added to whole blood at a concentration of 1×10^9 /mL and their interactions with leukocytes assayed by flow cytometry. Many monocytes acquired GPIb α within 5 min, but neutrophils or lymphocytes did not (Figure 3C-D). The proportion of monocytes acquiring GPIb α slowly increased thereafter, while intensity of GPIb α staining increased steadily (Figure 3C-D). Interestingly, we found that most of the GPIb α signal detected by flow cytometry was intracellular (~80%) on both monocyte subsets (*Online Supplementary Figure S8A-B*) and in agreement with the confocal imaging data described in Figure 2C. We also analysed the adhesion of PEV labelled with the lipophilic dye PKH67 to exclude antibody-mediated interaction of PEV with monocytes. PKH67 labelled all of the PEV in the activated-platelet supernatant (Figure 3E). The dynamics of PEV binding to monocytes, neutrophils or lymphocytes (Figure 3F) was similar to that for the antibody-labelled PEV (Figure 3C).

The mechanistic basis for the preferential accumulation of GPIb α on monocytes was investigated using adhesion-blocking reagents. Inclusion of a function-neutralising

anti-P-selectin antibody inhibited GPIb α accumulation on both monocytes and neutrophils, strongly implicating this platelet receptor in heterotypic adhesion with the two cells (Figure 4A and *Online Supplementary Figure S9A-B*). We measured the density of the P-selectin counter receptor P-selectin Glycoprotein Ligand 1 (PSGL-1) on blood leukocytes because the efficiency of GPIb α accumulation might reflect the surface density of this molecule. Figure 4B shows that there is substantially more PSGL-1 on monocytes than neutrophils, which could account for the differential levels of GPIb α accumulating on these cells. However, T cells, which did not accumulate GPIb α , also possessed abundant PSGL-1. Thus additional and cell specific adhesive interactions may be

required to stabilise P-selectin mediated adhesion under shear. Using a panel of function-neutralising antibodies against known platelet and leukocyte adhesion molecules, we could find no contribution to heterotypic aggregate formation from CD31, ICAM-2 or β 2-integrins (Figure 4C-E). However, an anti-phosphatidylserine (PS) antibody significantly reduced GPIb α accumulation on both neutrophils and monocytes (Figure 4C-E). The function of PS as a stabilising interactant is concordant with its documented patterns of interaction with monocytes and neutrophils, while its potential lack of interaction with T cells would account for the lack of GPIb α accumulation on these cells.

An important question was whether monocyte activa-

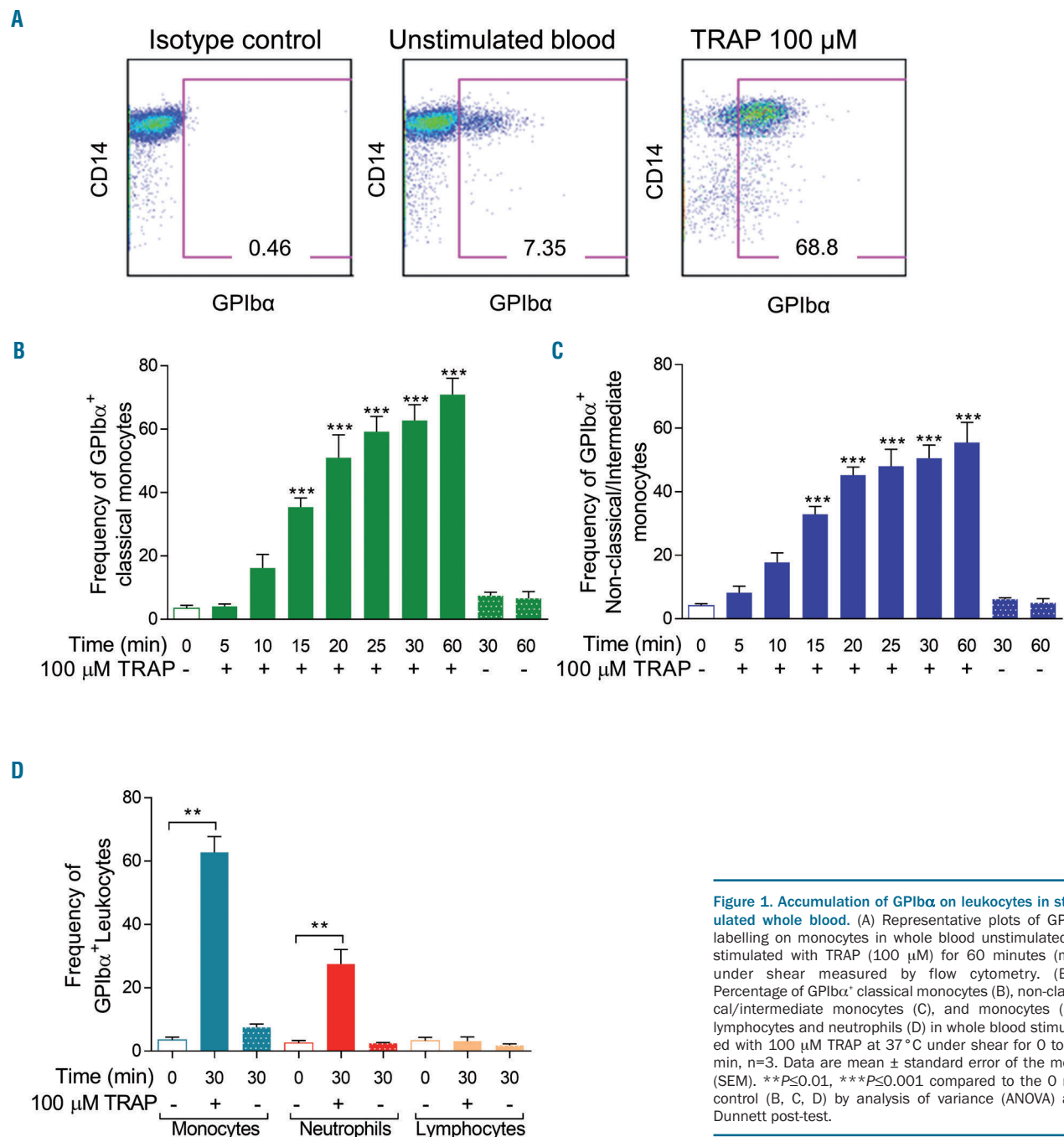


Figure 1. Accumulation of GPIb α on leukocytes in stimulated whole blood. (A) Representative plots of GPIb α labelling on monocytes in whole blood unstimulated or stimulated with TRAP (100 μ M) for 60 minutes (min) under shear measured by flow cytometry. (B-D) Percentage of GPIb α ⁺ classical monocytes (B), non-classical/intermediate monocytes (C), and monocytes (all), lymphocytes and neutrophils (D) in whole blood stimulated with 100 μ M TRAP at 37 °C under shear for 0 to 60 min, n=3. Data are mean \pm standard error of the mean (SEM). ** P ≤0.01, *** P ≤0.001 compared to the 0 min control (B, C, D) by analysis of variance (ANOVA) and Dunnett post-test.

tion contributed to GPIb α accumulation. For these studies, we used CRP-XL to stimulate whole blood, as this agonist does not directly activate monocytes and would thus allow analysis of whether secondary activation of monocytes downstream of platelet activation was prerequisite for PEV adhesion. We assessed the expression of the activation marker α M β 2-integrin (CD11b/CD18) on monocytes 30 min after the addition of CRP-XL to whole blood. There was some increase in both integrin subunits CD11b and CD18 (*Online Supplementary Figure S9C-D*), however, this was inconsistent and monocyte subset specific. When a function neutralising antibody against CD18 was included in the assay it had no effect on GPIb α accumulation (Figure 4C-E), indicating that monocyte activation was not required for PEV adhesion.

Adopted GPIb α is a functional adhesion molecule supporting monocyte rolling on von Willebrand Factor

As GPIb α is known to mediate binding of platelets from flowing blood to von Willebrand Factor (VWF), we tested whether VWF could also recruit PEV-treated monocytes (Figure 5A-E). Monocytes lacking GPIb α showed low levels of adhesion when perfused across immobilised human VWF (Figure 5B, E). However, acquisition of PEV-derived-GPIb α supported capture and rolling (66.8 \pm 4.1% of adherent cells rolling) of monocytes on VWF (Figure 5C, E). Importantly, the adhesion of PEV-treated mono-

cytes was inhibited by a function-neutralising antibody against GPIb α (Figure 5D, E).

Monocytes bearing GPIb α bind to EC in a model of vascular inflammation

Transforming growth factor beta-1 (TGF- β 1) promotes the expression of a matrix of VWF on the surface of EC which recruits platelets from flowing blood, which in turn function as adhesive bridges for the preferential recruitment of monocytes to EC *in vitro* and *in vivo*.¹⁵ Here we used this model to determine whether PEV-derived GPIb α could support monocyte adhesion directly to stimulated endothelium. A low level of monocyte adhesion to TGF- β 1-stimulated EC was observed without PEV (Figure 6A, D). However, PEV-treated monocytes adhered in significantly higher numbers, an adhesive interaction blocked by a GPIb α blocking antibody (Figure 6B-D). As previously observed, recruited monocytes did not roll on the EC. Thus 6.1 \pm 0.9% of adherent cells were observed rolling, with the remaining 93.9% becoming activated and stably adherent. Interestingly, the acquisition of PEV increased the efficiency with which monocytes transmigrated across the EC monolayer (Figure 6E). We could attribute this increase in PEV-treated monocytes recruitment to PEV rather than soluble factors such as chemokines, as supernatants generated from PEV filtered using a 10 KDa size filter (to remove

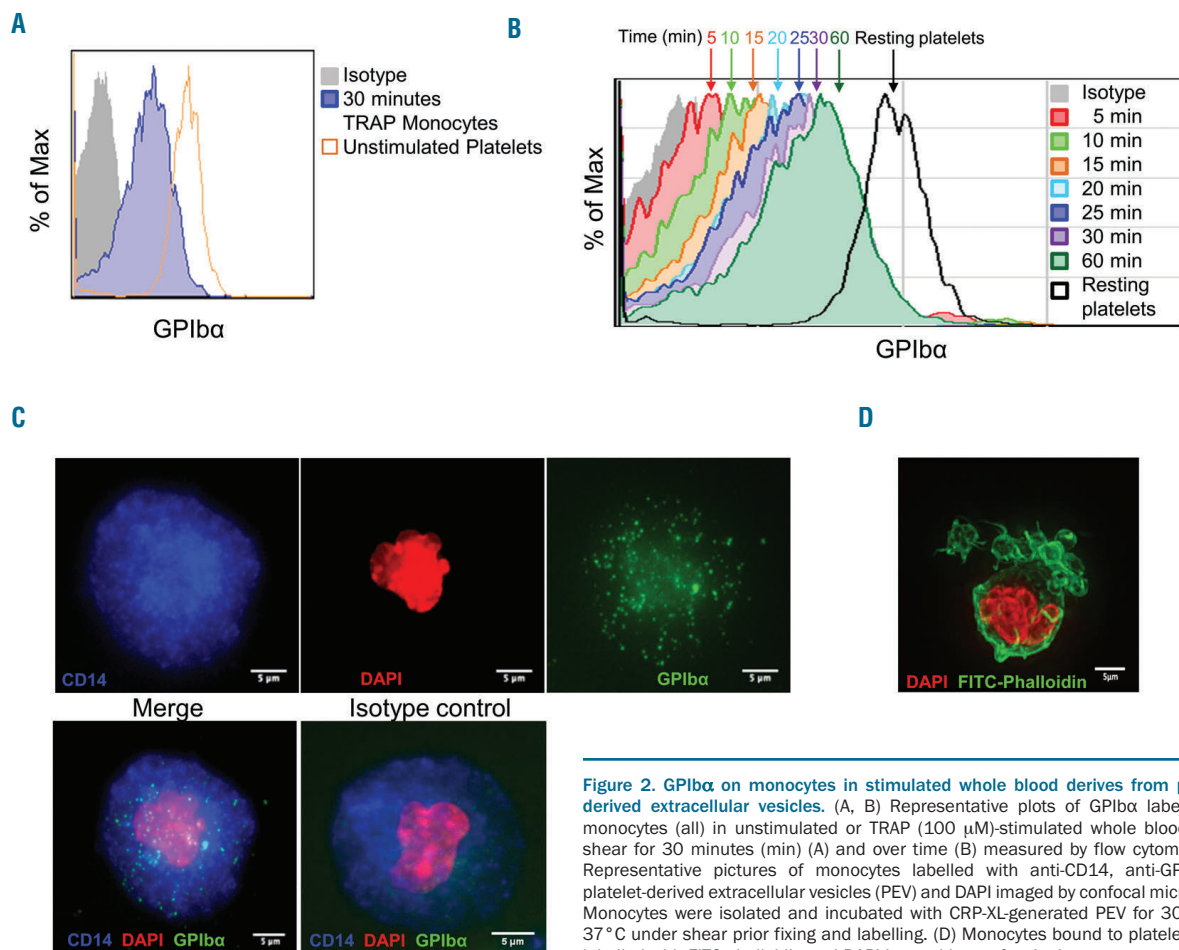


Figure 2. GPIb α on monocytes in stimulated whole blood derives from platelet-derived extracellular vesicles. (A, B) Representative plots of GPIb α labelling on monocytes (all) in unstimulated or TRAP (100 μ M)-stimulated whole blood under shear for 30 minutes (min) (A) and over time (B) measured by flow cytometry. (C) Representative pictures of monocytes labelled with anti-CD14, anti-GPIb α for platelet-derived extracellular vesicles (PEV) and DAPI imaged by confocal microscopy. Monocytes were isolated and incubated with CRP-XL-generated PEV for 30 min at 37 °C under shear prior fixing and labelling. (D) Monocytes bound to platelets, both labelled with FITC-phalloidin and DAPI imaged by confocal microscopy.

vesicles) did not induce monocyte adhesion and transmigration (Figure 6D, E).

PEV-treated murine monocytes bearing GPIIb α can be generated and recruited in mice

Prior to moving to *in vivo* assays of monocyte recruitment, we determined whether murine PEV derived-GPIIb α could accumulate on murine monocytes. Using the *ex vivo* whole blood assay under shear, we observed a high proportion of murine monocytes rapidly accumulated GPIIb α and CD41 after addition of ADP to the blood (Figure 7A and Online Supplementary Figure S10A-C). To examine

monocytes/PEV aggregate formation *in vivo* we induced pulmonary inflammation by instillation of air pollution particles into the lungs. A significant increase in the number of monocytes bearing GPIIb α and CD41 (α IIb-integrin) was observed in animals exposed to air pollution particles, but not vehicle control (PBS) (Figure 7B-C). Importantly, and in concordance with human studies, GPIIb α and CD41 intensities of expression was below the level on individual platelets (Online Supplementary Figure S10D), demonstrating that monocytes bind PEV in this model.

Using an intravital preparation of the TGF- β 1-stimulated, mouse cremaster muscle to observe monocyte interactions with the microvasculature in real time, we tracked

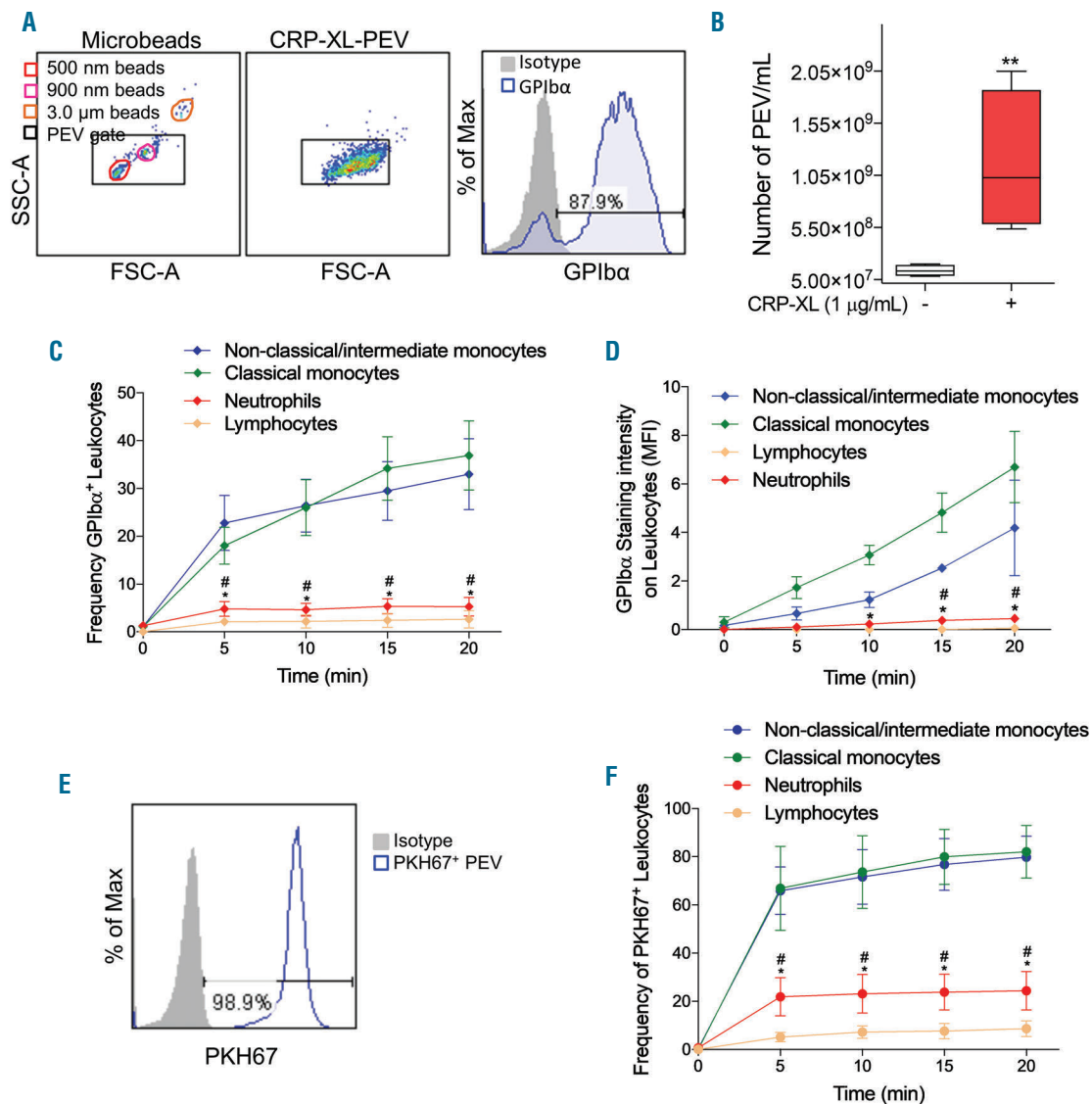


Figure 3. Rapid and specific binding of platelet-derived extracellular vesicles to monocytes. (A) Platelet-derived extracellular vesicles (PEV) gate was determined using microbeads to select events under 900 nm in size (left dot plots). Representative plot of GPIIb α intensity on PEV generated by stimulation of platelets with 1 μ g/mL CRP-XL for 30 minutes (min) analysed by flow cytometry (right histogram). (B) Concentration of PEV generated by stimulation of platelets with 1 μ g/mL CRP-XL for 30 min, n=4. (C, D) Percentage (C) and median fluorescent intensity (MFI) (D) of GPIIb α leukocytes in unstimulated whole blood supplemented with CRP_XL (1 μ g/mL)-generated-PEV at 37 °C under shear determined by flow cytometry, n=4. (E) Representative plot of PEV generated by stimulation of platelets with 1 μ g/mL CRP-XL for 30 min labelled with PKH67. (F) Percentage of PKH67⁺ leukocytes in unstimulated whole blood supplemented with CRP_XL (1 μ g/mL) generated PEV at 37 °C under shear determined by flow cytometry, n=4. Data are mean \pm standard error of the mean (SEM). ** $P \leq 0.01$ by (B) unpaired t-test. * or * $P \leq 0.01$ (C, D, F) by repeated measures two-way ANOVA followed by Bonferroni post-test for neutrophils and lymphocytes compared to * classical monocytes and # to non-classical/Intermediate monocytes.

human mouse PEV-treated monocytes (Online Supplementary Figure S10E-F). We used the hIL4R/GPIb α -Tg mouse which expressed human IL-4 receptor under the GPIb α promoter. This allows the animals to be rendered thrombocytopenic by injection of an antibody against hIL4R. Adoptively transferred WT platelets or PEV are however retained within the circulation. Using mice depleted of endogenous platelets using an anti-hIL4R antibody, we observed higher numbers of adoptively transferred WT PEV-treated monocytes rolling on the microvasculature compared to untreated monocytes; the number was significantly reduced by a GPIb α blocking antibody (Figure 7D-G). Detailed analysis revealed two populations of rolling cells: those exhibiting stable rolling (interactions >300 ms) with a velocity of 241 ± 82 $\mu\text{m/s}$ (Figure 7D, F); those exhibiting transient rolling (interactions <300 ms) with a velocity of 478 ± 65 $\mu\text{m/s}$ (Figure 7D, G). We also infused human monocytes

into *ApoE*^{-/-} mice that had been on a western diet for six weeks and observed the carotid artery by intravital microscopy. Murine PEV-treated monocytes adhered to the artery wall with significantly greater efficiency than untreated monocytes (Figure 7H). In this environment a mixture of adhesive behaviors was observed with stationary adhesion, stable rolling and transient rolling adhesion evident (Figure 7H).

Monocytes with platelet markers appear within 1 hour of severe trauma and are rapidly cleared from the circulation

We investigated whether rapid production and binding of extracellular vesicles to monocytes could be detected following an acute event such as traumatic injury. In the Golden Hour study blood samples in the pre-hospital setting (mean time to blood sampling = 43 min) were acquired from traumatically-injured patients (injury sever-

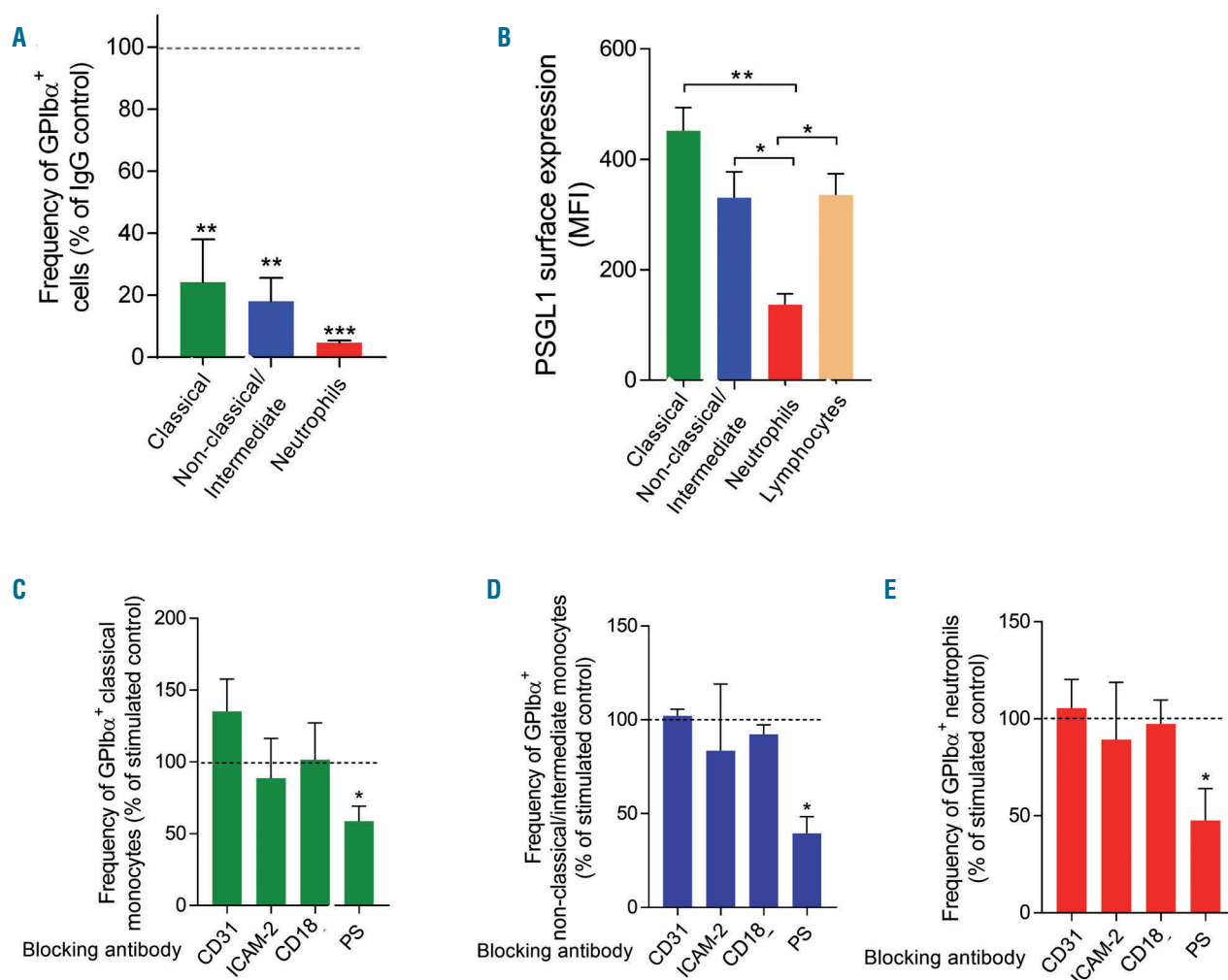


Figure 4. Blocking of GPIb α ⁺ platelet-derived extracellular vesicles to leukocytes. (A) Binding of platelet-derived extracellular vesicles (PEV) on classical monocytes, non-classical/intermediate monocytes and neutrophils with blockade of P-selectin in TRAP (100 μM)-stimulated whole blood for 30 minutes (min) under shear, $n=3$. (B) Surface expression (MFI: median fluorescence intensity) of PSGL1 (P-selectin ligand) on monocyte subsets, neutrophils and lymphocytes determined by flow cytometry, $n=3$. (C-E) Binding of PEV on classical monocytes (C), non-classical/intermediate monocytes (D) and neutrophils (E) with blockade of CD31, ICAM-2, CD18 ($\beta 2$) and Phosphatidylserine (PS) in TRAP (100 μM)-stimulated whole blood for 30 min under shear, $n=3-5$. Data are mean \pm standard error of the mean (SEM). * $P \leq 0.05$, ** $P \leq 0.01$ compared the normalised IgG control (A) by analysis of variance (ANOVA) and Dunnett post-test or Bonferroni post-test (B) and one sample t-test to 100% of TRAP control (C-E).

ity score [ISS] >8). Analysis by flow cytometry showed acquisition of CD41 by circulating leukocytes with preferential binding to monocytes (Figure 8A-B and *Online Supplementary Figure S11A-B*). The CD41 measured on monocytes was likely derived from PEV, as the intensity of fluorescent staining at 4 hours post trauma ($2,226 \pm 474$) was substantially below that of a single platelet ($13,702 \pm 964$) (*Online Supplementary Figure S11B*). Both the number of CD41⁺ monocytes and the intensity of staining for CD41 on them (MFI), correlated significantly with the severity of trauma (Figure 8C-D). Lastly, there was a marked loss of CD41⁺ monocytes from the blood within 4 to 12 hours, which was sustained for up to 72 hours (Figure 8E) and a decrease in circulating platelet counts which reflects platelet activation and PEV generation (*Online Supplementary Figure S11C*).

Discussion

We have defined a new thrombo-inflammatory route of monocyte recruitment *via* an adhesion molecule transferred from platelets. Recruitment is reliant upon platelet-derived GPIb α , which allows monocyte capture by VWF exposed on the vessel wall. Previous studies have indicated that platelet-derived chemokines can then induce arrest and migration.⁴⁵ *Online Supplementary Figure S12* summarises the steps we propose in this thrombo-inflammatory cascade. Importantly, the cascade may diverge from the normal pathways of leukocyte trafficking in a manner that could contribute to disease, as plasma borne PEV preferentially deliver functional GPIb α to the monocyte surface. Transfer of GPIb α can support adhesion of monocytes *in vitro* and *in vivo*, in human and murine models of

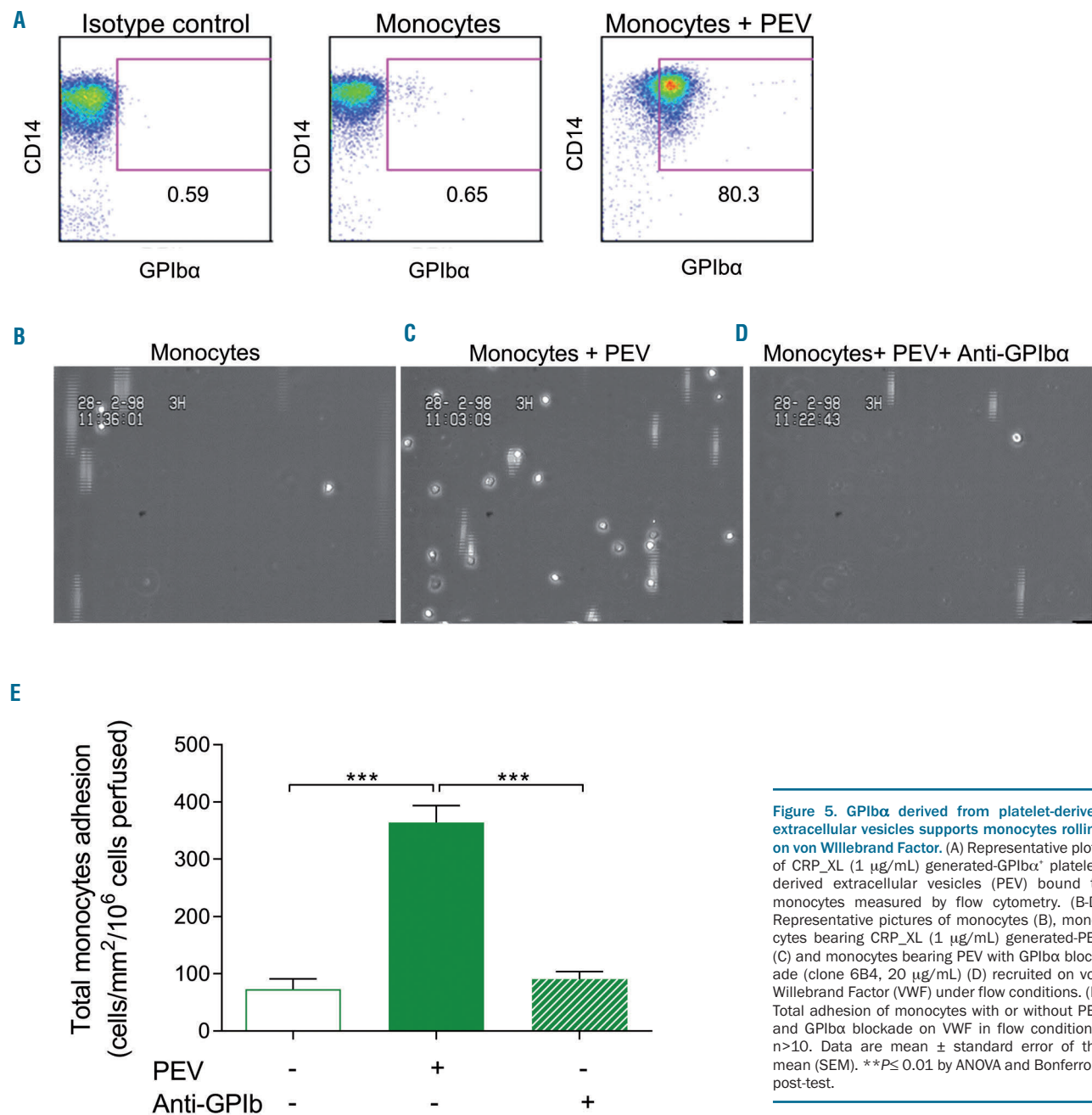


Figure 5. GPIb α derived from platelet-derived extracellular vesicles supports monocytes rolling on von Willebrand Factor. (A) Representative plots of CRP_XL (1 μ g/mL) generated-GPIb α ⁺ platelet-derived extracellular vesicles (PEV) bound to monocytes measured by flow cytometry. (B-D) Representative pictures of monocytes (B), monocytes bearing CRP_XL (1 μ g/mL) generated-PEV (C) and monocytes bearing PEV with GPIb α blockade (clone 6B4, 20 μ g/mL) (D) recruited on von Willebrand Factor (VWF) under flow conditions. (E) Total adhesion of monocytes with or without PEV and GPIb α blockade on VWF in flow conditions, $n > 10$. Data are mean \pm standard error of the mean (SEM). *** $P \leq 0.01$ by ANOVA and Bonferroni post-test.

vascular inflammation. This process means that circulating monocytes may be recruited to the vessel wall through a pathway outside of the tightly regulated physiological inflammatory system. We believe that such monocyte recruitment may be particularly relevant in the dysregulated inflammatory responses seen in chronic inflammatory disease, which leads to tissue damage and loss of function (such as atherosclerosis and rheumatoid arthritis). In addition, it may be important in inflammation associated with severe trauma, where the drivers of inflammation are substantial and acute tissue damage, and extensive activation of the coagulation and haemostatic pathway. However, we believe that during acute responses initiated by inflammatory cytokines in a coordinated and controlled manner, and where timely and comprehensive resolution is the norm, platelet-mediated pathways of leukocyte trafficking are likely to be of lesser importance.

Other studies show that whole platelets can bind leukocytes, a process dependent upon platelet and/or leukocyte activation and linked to pathological conditions.⁴⁶ Moreover, if PEV are mixed with isolated monocytes they are able to activate the leukocytes so that they show enhanced levels of recruitment to EC *in vitro*, although

direct binding between PEV and leukocytes was not demonstrated in that study.⁴⁷ Here, we show that monocytes preferentially accumulate PEV rather than whole platelets through an adhesive pathway reliant upon P-selectin. In the context of leukocyte recruitment to vascular EC, P-selectin supports a distinct form of rolling adhesion which is based on the transient nature of the bonds formed with PSGL-1 under conditions of shear.⁴⁸ Here we propose that the P-selectin-PSGL-1 mediated interactions between PEV and leukocytes are also transient under the shear conditions of our assay and in flowing blood *in vivo*. However, on monocytes and neutrophils, PS in the PEV membrane acts to stabilise heterotypic adhesion upon interaction with membrane receptors on the leukocytes. In the case of T lymphocytes, which also possess abundant PSGL-1, the transient interactions formed with P-selectin under shear are not stabilised by PS, which has not been reported to bind T cells to our knowledge.

In fact, much of the data on heterotypic aggregate formation in human blood does not discriminate between platelets and PEV binding to leukocytes, and it is unclear which is being assessed. Studies that do report platelet

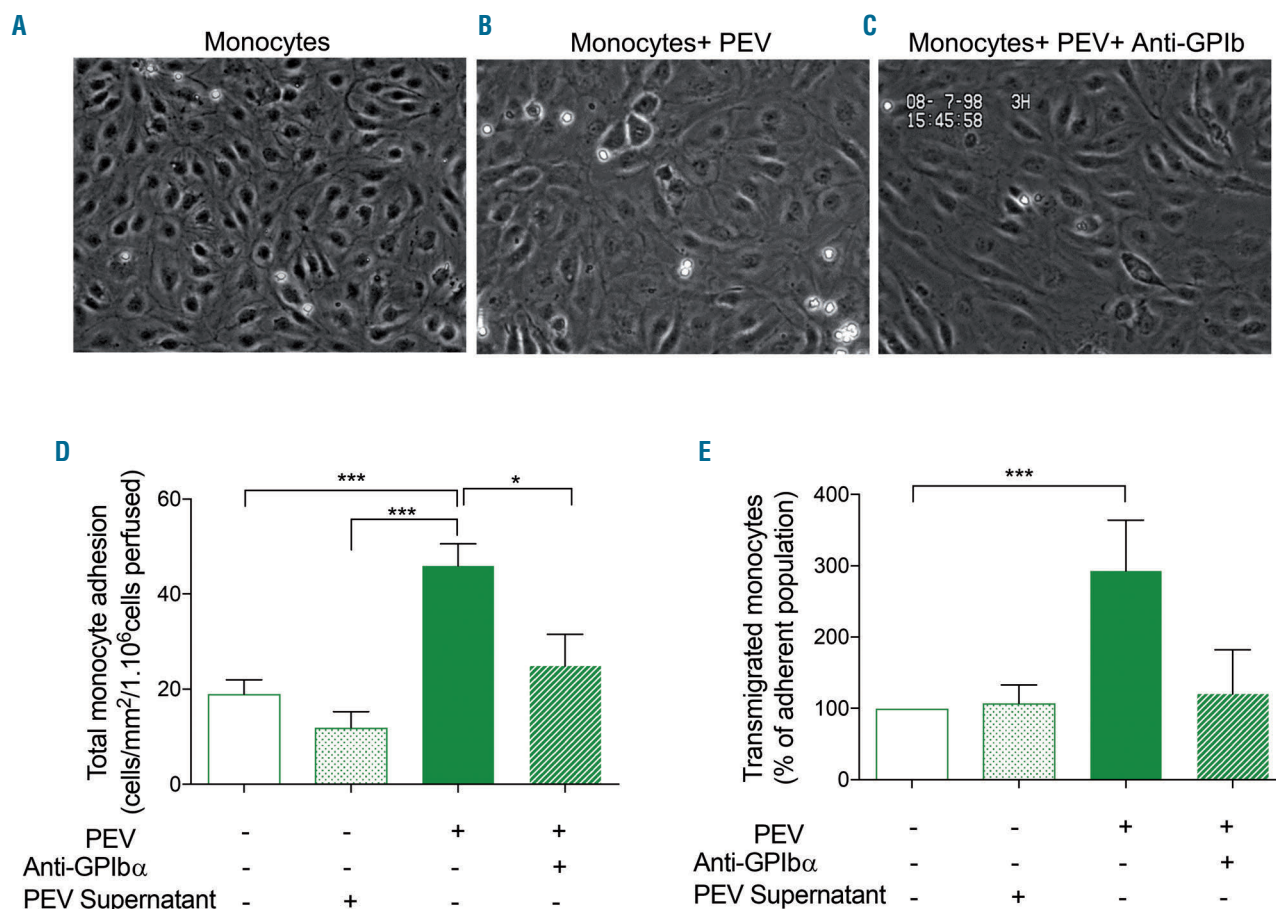


Figure 6. GPIIb/IIIa derived from platelet-derived extracellular vesicles supports monocytes recruitment on TGF- β 1 stimulated endothelial cells. (A-C) Representative pictures of monocytes (A), monocytes bearing CRP_XL (1 μ g/mL) generated-PEV (B) and monocytes bearing PEV with GPIIb/IIIa blockade (clone 6B4, 20 μ g/mL) (C), adhered on TGF- β 1 (10 ng/mL) stimulated EC in flow conditions. (D, E) Total adhesion (D) and transmigration (E) of monocytes with or without PEV, GPIIb/IIIa blockade and filtered PEV through a 10 kDa filters to remove PEV and leave potential soluble factors on TGF- β 1 stimulated EC in flow conditions, n=3-5. Data are mean \pm standard error of the mean (SEM). * $P \leq 0.05$, ** $P \leq 0.01$ by ANOVA and Bonferroni post-test.

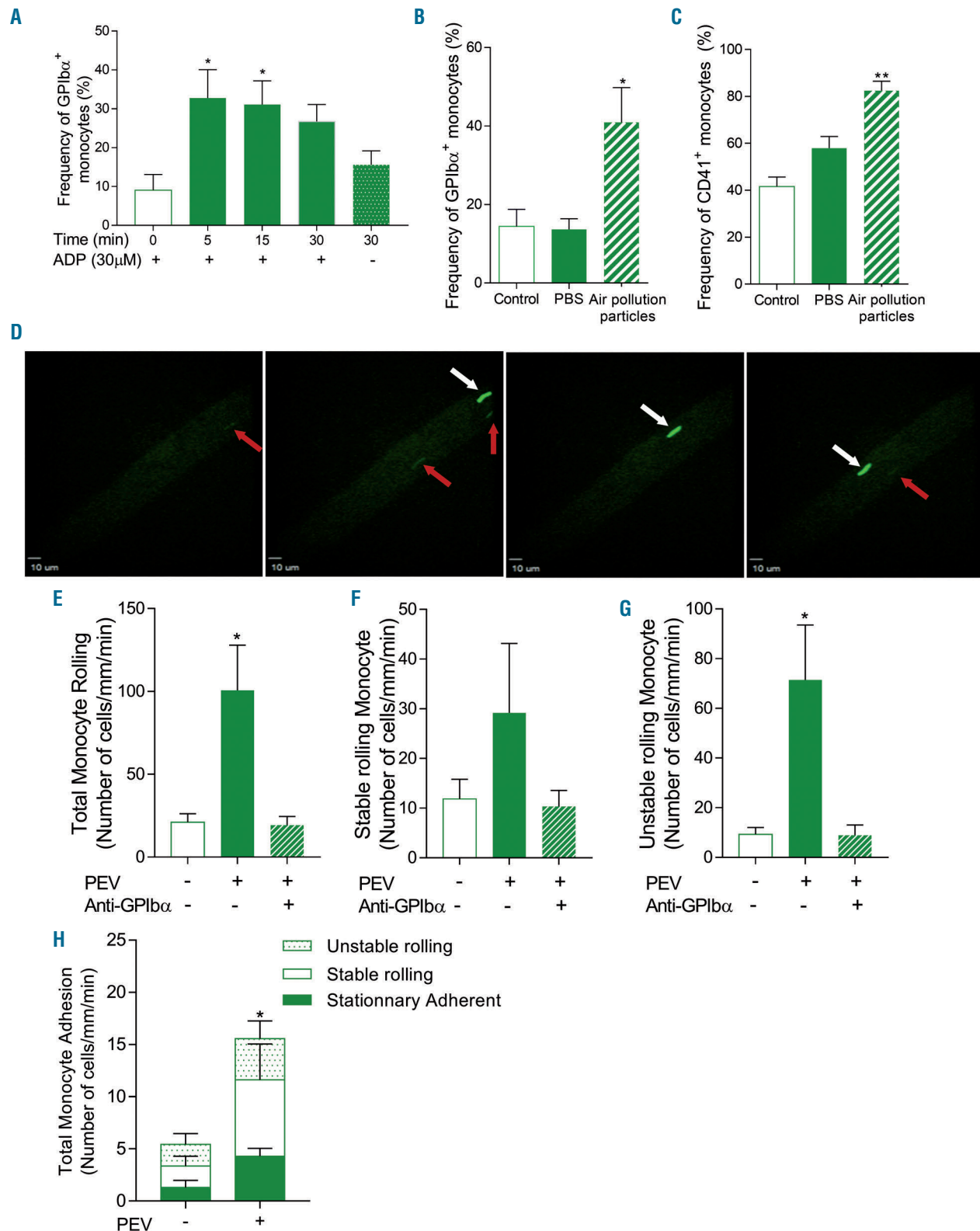


Figure 7. GPIIb/IIIa from platelet-derived extracellular vesicles mediates recruitment of monocytes *in vivo*. (A) Percentage of GPIIb/IIIa⁺ monocytes in whole murine blood stimulated with 30 μM ADP at 37 °C under shear determined by flow cytometry, n=4. (B) Percentage of GPIIb/IIIa⁺ and (C) CD41⁺ monocytes in murine blood 48 hours after intratracheal instillation of air pollution particles (400 μg) measured by flow cytometry, n=5-6. (D) Representative images of the recruitment of human monocytes bearing GPIIb/IIIa from mouse platelet-derived extracellular vesicles (PEV) to the TGF-β1 (80 μg/kg)-stimulated vasculature in the cremaster muscle using intravital microscopy. Red arrows point to unstable rolling monocytes, white arrows to stable rolling monocytes. Total rolling (E) stable rolling (F) and unstable rolling (G) of human monocytes bearing GPIIb/IIIa from mouse PEV with or without with GPIIb/IIIa blockade (clone Xia.B2, 50 μg/mL) adoptively transferred in the platelet depleted IL4R/GPIIb/IIIa-Tg mice measured by intravital microscopy of the TGFβ1-stimulated cremaster muscle, n=3. (H) Total adhesion and behaviors of human monocytes bearing mouse PEV adoptively transferred in western diet fed ApoE^{-/-} mice measured by intravital microscopy of the right carotid artery, n=3. Data are mean ± standard error of the mean (SEM). *P≤0.05, **P≤0.01 compared to the 0 minute (min) control (A) or control (B, C) by analysis of variance (ANOVA) and Dunnett post-test and by unpaired t-test, by two-way ANOVA (H).

binding are routinely performed *ex vivo* under the non-physiological condition of stasis *in vitro*, where the number of platelet-leukocyte aggregates formed is a direct function of the time of incubation.⁴⁹ Thus, patient blood may have a greater propensity to form aggregates with platelets under static conditions *ex vivo*, but this probably does not reflect the situation *in vivo*. Such aggregation may be a surrogate endpoint for the degree of platelet and/or leukocyte activation present in patient blood. In support of this, the patterns of PEV associated with circulating monocytes that we report here are in strong accord with a recent report from Fendl *et al.* who analysed the effects of pre-analytical blood handling (which included the imposition of shear) on the association of extracellular vesicles with leukocytes.⁵⁰

Interestingly, upon addition of purified PEV to whole blood, we observed rapid accumulation of GPIb α on monocytes, implying assimilation of PEV was extremely efficient. However, when a platelet activating agonist was added to whole blood the process was continuous and prolonged, leading to an incremental increase in GPIb α expression. The latter profile of accumulation of GPIb α likely reflects the dynamics of PEV formation and release

by platelets in whole blood, implying that the rate-limiting step in this thrombo-inflammatory pathway is not PEV-monocyte interaction, but rather the process of PEV release after platelet activation. In addition, accumulation of PEV was more prevalent in monocytes compared to neutrophils and lymphocytes. In a previous study, we observed different patterns of recruitment, migration and reverse migration *in vitro* between classical and non-classical/intermediate monocytes.⁴³ We characterised a novel process of crosstalk mediated by cytokines between the two subsets that allowed a balanced regulation of endothelial cell activation. Other studies have shown that changes in proportional representation of monocyte subsets in the circulation are associated with vascular diseases.^{51,52} However, in this study we observed no preferential binding of PEV between classical and non-classical/intermediate monocytes, which was consistent with similar levels of PSGL1 expression exhibited by all subsets.

GPIb α is an adhesion receptor mediating a specialised form of platelet recruitment during haemostasis. Bonds forming between GPIb α and VWF exhibit high on rates, meaning that adhesion can occur between rapidly flowing

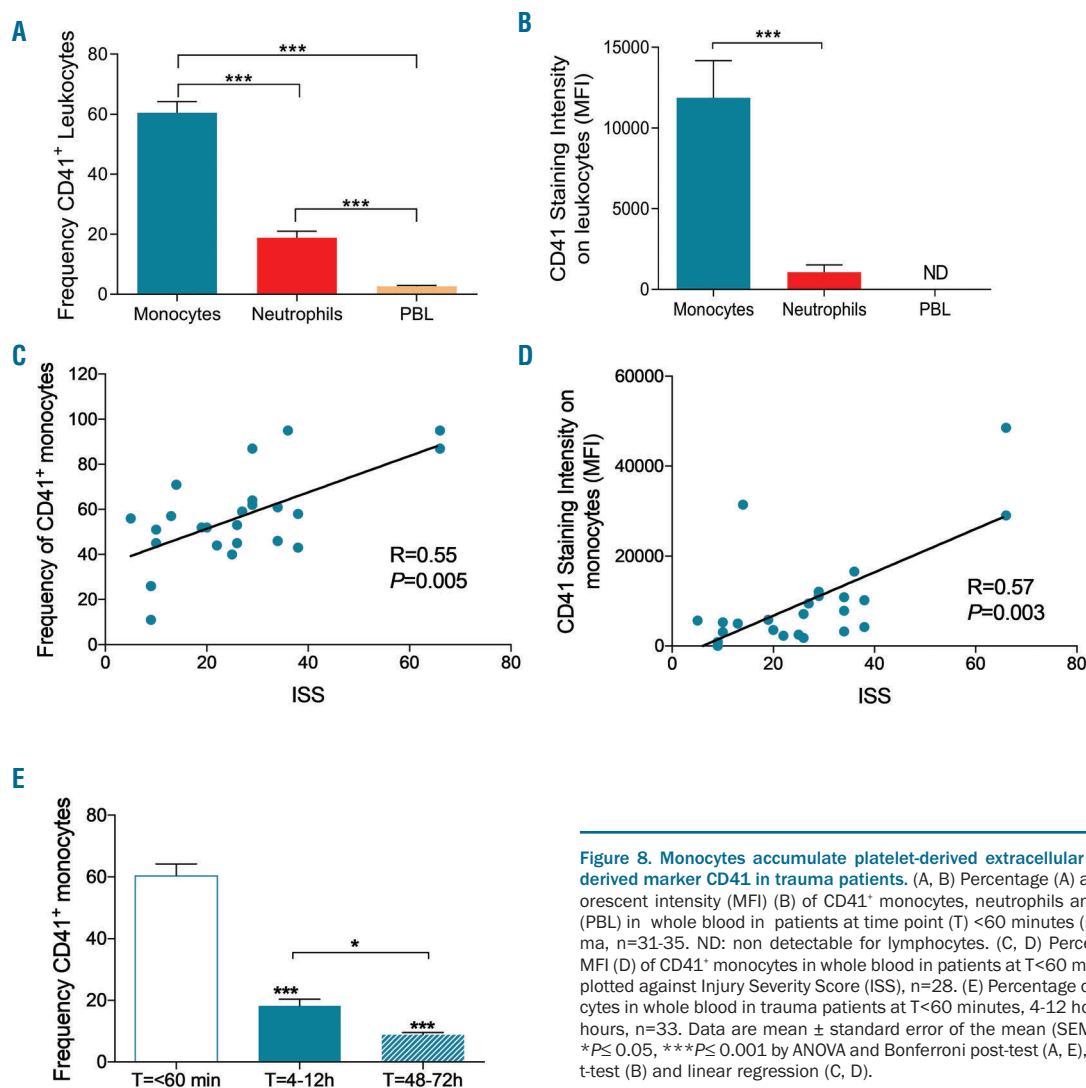


Figure 8. Monocytes accumulate platelet-derived extracellular vesicles (PEV) derived marker CD41 in trauma patients. (A, B) Percentage (A) and median fluorescent intensity (MFI) (B) of CD41⁺ monocytes, neutrophils and lymphocytes (PBL) in whole blood in patients at time point (T) <60 minutes (min) after trauma, n=31-35. ND: non detectable for lymphocytes. (C, D) Percentage (C) and MFI (D) of CD41⁺ monocytes in whole blood in patients at T<60 min after trauma plotted against Injury Severity Score (ISS), n=28. (E) Percentage of CD41⁺ monocytes in whole blood in trauma patients at T<60 minutes, 4-12 hours and 48-72 hours, n=33. Data are mean \pm standard error of the mean (SEM) (A, B and E). * $P \leq 0.05$, *** $P \leq 0.001$ by ANOVA and Bonferroni post-test (A, E), Mann Whitney t-test (B) and linear regression (C, D).

platelets and the substrate.^{15,53} However, these bonds also exhibit high off rates. Thus, under conditions of shear stress (*i.e.* blood flow) the rapid formation and dissolution of bonds supports rolling adhesion.^{15,53} We observed that monocytes bearing GPIb α also rolled on purified VWF. However, on EC bearing VWF, monocytes were rapidly activated, which is consistent with our previous observations on the activity of EC derived stimuli such as C-C chemokine ligand 2 (CCL2).¹⁵

Trogocytosis is the phenomenon by which lymphocytes extract surface molecules from antigen presenting cells through transfer of plasma membrane at the immunological synapse.⁵⁴ This process has been observed for T, B and NK cells and neutrophils^{54,55} and is a fast and efficient means of transferring molecules involved in the regulation of immune functions.⁵⁴ We cannot completely exclude that monocytes in whole blood do not bind whole platelets and acquire GPIb α and CD41 via a trogocytosis like process, although a synapse like structure has not been reported in this context to our knowledge. However, both trogocytosis and PEV accumulation by monocytes require activation dependent cytoskeletal rearrangement to achieve the transfer of membrane cargo that alters the function of the recipient cells. Thus, the processes may not be unrelated in their mechanisms of initiation and progression. However, trogocytosis does appear to be specific to the immunological synapse or related structures.^{54,56} Here however, we have shown that purified labelled PEV bind to isolated monocytes or monocytes in whole blood with the same dynamics as agonist stimulated systems. This clearly demonstrates that a trogocytosis like process is not required for accumulation of PEV once they have been generated by platelet activation (Figure 3). Our colleagues have also shown that PEV levels increase dramatically after trauma and thus are likely to be the source of GPIb α found on leukocytes in trauma patients.⁵⁷

Using intravital microscopy we observed GPIb α -dependent recruitment of PEV-treated monocytes to the vasculature. Interestingly, the short-lived adhesive interactions, here termed 'transient rolling' which did not result in prolonged monocyte localisation and activation at the vessel wall, have previously been shown to have physiological roles. Thus, under steady-state conditions (non-inflamed), circulating platelets expressing GPIb α are able to interact transiently with sinusoidal Kupffer cells in the liver *via* surface-expressed VWF, interactions which are important for host defence, as they facilitate uptake and disposal of bacteria by liver resident macrophages (Kupffer cells).⁵⁸ In the context of CVD, we showed that induction of pulmonary inflammation with pollution nanoparticles, a known risk factor for thrombo-inflammatory disease associated with atherosclerosis,⁵⁹ induced the formation of circulating monocyte-PEV aggregates. Moreover, such aggregates showed a significantly enhanced capacity to bind to the artery wall in the *ApoE*^{-/-} mouse after induction of disease by feeding a high fat western diet. Thus, we propose that the transfer of platelet cargo to monocytes by PEV can contribute to the progression of plaque formation by promoting the recruitment of inflammatory monocytes. It would be interesting to investigate the functional and phe-

notypical changes induced by binding and internalisation of PEV by monocytes. In this study we did not observe major changes in integrin expression as a marker of activation. However, we do not exclude changes in monocyte activation and/or function relevant to vascular disease over longer periods of interaction.

The paradigm discussed above may provide a novel thrombo-inflammatory mechanism for the continuous low levels of monocyte delivery in chronic inflammatory conditions such as atherosclerosis. However, our Golden Hour data suggest that acquisition of this pathway of monocyte recruitment could also lead to the clearance of monocytes from the blood during acute and severe trauma. Indeed, in this injured patient cohort, PEV counts increase in the circulation⁵⁷ and monocytes rapidly acquired CD41-derived from PEV (1 hour after trauma) which we believe led to their clearance from the circulation, as frequency of CD41⁺ monocyte numbers are lower 4 hours after trauma. This may be due to clearance by the reticulo-EC system, or alternatively by the expedited recruitment to damaged and inflamed tissues, or indeed a combination of both. Whatever the pathway of their removal from the circulation, we speculate that the rapid clearance of immune cells from the circulation may exacerbate cell turnover and result in immune suppression and the increased risk of septic complications. In addition, monocytes bearing pro-coagulant PEV could also contribute to the initiation and propagation of disseminated intravascular coagulation (DIC) which is a potential and serious complication of traumatic injury.²⁷

In conclusion, we believe that this new paradigm for leukocyte recruitment is an important step in understanding the contribution of platelets to thrombo-inflammatory pathology. By acquiring GPIb α in the circulation, monocytes may be provided with a means of interacting with the vessel wall, which is ordinarily restricted to platelets during haemostasis. In chronic diseases such as atherosclerosis, this process may occur with a low frequency over protracted periods of time. Nevertheless, the dynamic nature of PEV-monocyte interaction demonstrated in this study implies that such routes of thrombo-inflammation may be major contributors to pathology.

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